

**K955352 SUCTION CAUTERY PROBE**Apr 5, 1996  
136 days to decisionK955352 · Product code: **GEI** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k955352/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	Nov 21, 1995
Decision date	Apr 5, 1996
Days to decision	136 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

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Company	<b>Mectra Labs, Inc.</b>
Location	Bloomfield, IN, US
Contact	CHARLES E ALLGOOD
510(k) history	9 submissions · 9 cleared · 1992-2012

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k955352/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at [accessdata.fda.gov](http://accessdata.fda.gov). - Generated June 14, 2026